

us-09-832-658-7.n2p.rag

XX	KW	cytostatic; virucide; hepatotropin; antianaphylactic; treatment; fibrosis;
PR	KW	multiple sclerosis; inflammatory disease; autoimmune disease; cancer;
IR	KW	hepatitis; viral infection; neovascularisation; IFN beta 1a; helix A;
XX	OS	Homo sapiens.
PA	XX	location/Qualifiers
PA	XX	1..25
PA	KEY	/note- Helix_A
PA	PT	Misc-difference 2
PA	PT	/note- "Wild type Ser is substituted by Ala"
PA	PT	Misc-difference 4
PA	PT	/note- "Wild type Asn is substituted by Ala"
PA	PT	Misc-difference 5
PA	PT	/note- "Wild type Ieu is substituted by Ala"
PA	PT	Misc-difference 8
PA	PT	/note- "Wild type The is substituted by Ala"
PA	PT	Misc-difference 11
PA	PT	/note- "Wild type Ara is substituted by Ala"
PA	PT	26..53
PA	Domain	/label- AB_loop
PA	PT	54..73
PA	PT	/label- Helix_B
PA	PT	74..100
PA	PT	/label- Helix_C
PA	PT	101..119
PA	PT	/label- CD_loop
PA	PT	120..134
PA	PT	/label- DE_loop
PA	PT	135..140
PA	PT	/label- Helix_E
PA	PT	141..166
PA	PT	/label- Helix_F
PA	PT	PR:200022477-A2.
PA	XX	
PA	TC	27 APR-2000.
PA	XX	
PA	PR	15-OCT-1999; 9900-US24200.
PA	XX	
PA	PR	16-OCT-1998; 98US-0144491.
PA	PR	16-FEB-1999; 99US-0120237.
PA	XX	
PA	(HIOJ) BIOPRO INC.	
PA	XX	
PA	KEY	Whittier, A.; Forster, T.; Bruckelmaier, M.; Hochman, P;
PA	PT	WIT; 2000-1396547-29.
PA	PT	Fusion Proteins comprising interferon-beta-1a useful for inhibiting
PA	PT	angiogenesis -
PA	XX	Example 1; Page : 82pp; English.
PS	XX	The patient discloses fusion proteins comprising glycosylated
PS	XX	interferon-beta (IFN beta) especially IFN-beta-1a, linked groups and
PS	XX	non-IFN-beta proteins especially an immunodominant (Ig) protein. The
PS	XX	fusion protein is useful for inhibiting angiogenesis in a patient.
PS	XX	It may also be used to treat multiple sclerosis, tibiosis, inflammation
PS	XX	and autoimmune diseases, cancers, hepatitis and viral infections.
PS	XX	The present sequence is a human interferon-beta alanine substituted mutant A1, substituted by methionine at position 14, so 14-Met-15-Phe-16-Tyr-KWY72872.
PS	XX	a human interferon-beta alanine substituted mutant A1, substituted by
PS	XX	substituting residues in the helix A of wild type sequence.
PS	XX	The A1/Ser substituted mutants of IFN beta having substitutions in
PS	XX	helices A, B, C, D or E, or loops A, C or D were assessed for
PS	XX	receptor binding and functional activities. The A1 mutant shows antiviral and
PS	XX	anti-proliferative activities which are disproportionately low with
PS	XX	respect to receptor binding, compared to wild-type IFN-beta-1a. The
PS	XX	mutant can be used to produce IFN-beta fusion proteins
PS	XX	Note: The present sequence is not shown in the specification but is
PS	CC	derived from wild type human IFN-beta sequence found in page 36

an amino acid residue constituting an attachment group for a first non-polypeptide group and subjecting the modified peptide to conjugation with the non-polypeptide group. The conjugate and a cell culture expressing the mutated polypeptides are useful in the treatment of diseases, especially multiple sclerosis, and for treating mammals having mutated proteins may be used for gene therapy. The DNA and proteins can also be used to treat viral infections (e.g. viral hepatitis), cancer (e.g. breast cancer), inflammation, crohn's disease, acute myeloid leukaemia, Hodgkin's disease and ulcerative colitis and for immunomodulation.

Note: The present sequence is not shown in the specification but is derived from the human interferon beta sequence given in AUU0038.

XX sequence 166 AA;

XX

alignments_scores:
Quality: 90.00 Length: 22
Ratio: 5.000 Gaps: 0
Percent Similarity: 81.818 Percent Identity: 77.273

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Quality: 90.00 Length: 22
Ratio: 5.000 Gaps: 0
Percent Similarity: 81.818 Percent Identity: 77.273

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Quality: 90.00 Length: 22
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Quality: 90.00 Length: 22
Ratio: 5.000 Gaps: 0
Percent Similarity: 81.818 Percent Identity: 77.273

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alignments_scores:
Quality: 90.00 Length: 22
Ratio: 5.000 Gaps: 0
Percent Similarity: 81.818 Percent Identity: 77.273

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modified proteins may be used for gene therapy. The DNA and proteins can also be used to treat viral infections (e.g. viral hepatitis), cancer (e.g. breast cancer), inflammation (e.g. rheumatoid arthritis, acute myeloid leukaemia, psoriasis), a disease and therapeutic agents and therapeutic drugs.

Note: The first set of figures is not shown in the specification but is derived from the human interferon beta sequence given in AAN0038.

Sequence: 31 AA:

Alignment scores: Quality: 90.00 Length: 22

Similarity: 5.00 Gap: 0

Percent similarity: 81.81 Percent identity: 77.273

Alignment block: XX

Sequence: 31 AA:

Alignment scores: Quality: 89.00 Length: 22

Similarity: 4.944 Gap: 0

Percent similarity: 81.818 Percent identity: 77.273

Alignment block: XX

Sequence: 31 AA:

Alignment scores: Quality: 89.00 Length: 22

Similarity: 4.944 Gap: 0

Percent similarity: 81.818 Percent identity: 77.273

Alignment block: XX

Sequence: 31 AA:

Alignment scores: Quality: 89.00 Length: 22

Similarity: 4.944 Gap: 0

Percent similarity: 81.818 Percent identity: 77.273

Alignment block: XX

Sequence: 31 AA:

Alignment scores: Quality: 89.00 Length: 22

Similarity: 4.944 Gap: 0

Percent similarity: 81.818 Percent identity: 77.273

Alignment block: XX

Sequence: 31 AA:

Alignment scores: Quality: 89.00 Length: 22

Similarity: 4.944 Gap: 0

Percent similarity: 81.818 Percent identity: 77.273

Alignment block: XX

Sequence: 31 AA:

Alignment scores: Quality: 89.00 Length: 22

Similarity: 4.944 Gap: 0

Percent similarity: 81.818 Percent identity: 77.273

Alignment block: XX

Sequence: 31 AA:

Alignment scores: Quality: 89.00 Length: 22

Similarity: 4.944 Gap: 0

Percent similarity: 81.818 Percent identity: 77.273

Alignment block: XX

Sequence: 31 AA:

Alignment scores: Quality: 89.00 Length: 22

Similarity: 4.944 Gap: 0

Percent similarity: 81.818 Percent identity: 77.273

Alignment block: XX

Sequence: 31 AA:

CC to the 5' terminal of human fibroblast IFN, the entire sequence of mature human fibroblast interferon encoded by AAN0052 (see also AAPI051). Another recombinant IFN which was reported different from AAN0049 in that the codon for the sixth amino acid had a "silent" base change (TAC to TAT), indicating the existence of genetic polymorphism (see AAPI052/N0051).

CC Sequence: US-09-832-658-7 x AAF10052 . . .

CC Alignment scores: Quality: 89.00 Length: 22

CC Similarity: 4.944 Gap: 0

CC Percent similarity: 81.818 Percent identity: 77.273

CC Alignment block: 7.1

CC Sequence: 832-658-7 x AAF10052 . . .

CC Alignment scores: Quality: 89.00 Length: 22

CC Similarity: 4.944 Gap: 0

CC Percent similarity: 81.818 Percent identity: 77.273

CC Alignment block: 7.1

CC Sequence: 832-658-7 x AAF10052 . . .

CC Alignment scores: Quality: 89.00 Length: 22

CC Similarity: 4.944 Gap: 0

CC Percent similarity: 81.818 Percent identity: 77.273

CC Alignment block: 7.1

CC Sequence: 832-658-7 x AAF10052 . . .

CC Alignment scores: Quality: 89.00 Length: 22

CC Similarity: 4.944 Gap: 0

CC Percent similarity: 81.818 Percent identity: 77.273

CC Alignment block: 7.1

CC Sequence: 832-658-7 x AAF10052 . . .

CC Alignment scores: Quality: 89.00 Length: 22

CC Similarity: 4.944 Gap: 0

CC Percent similarity: 81.818 Percent identity: 77.273

CC Alignment block: 7.1

CC Sequence: 832-658-7 x AAF10052 . . .

CC Alignment scores: Quality: 89.00 Length: 22

CC Similarity: 4.944 Gap: 0

CC Percent similarity: 81.818 Percent identity: 77.273

CC Alignment block: 7.1

CC Sequence: 832-658-7 x AAF10052 . . .

CC Alignment scores: Quality: 89.00 Length: 22

CC Similarity: 4.944 Gap: 0

CC Percent similarity: 81.818 Percent identity: 77.273

CC Alignment block: 7.1

CC Sequence: 832-658-7 x AAF10052 . . .

CC Alignment scores: Quality: 89.00 Length: 22

CC Similarity: 4.944 Gap: 0

CC Percent similarity: 81.818 Percent identity: 77.273

CC Alignment block: 7.1

CC Sequence: 832-658-7 x AAF10052 . . .

PT It is understood that the probability of homologicity between human IFN variants (IFN- α and IFN- β) and human fibroblast IFN is higher than that between human fibroblast IFN and human lymphoblast IFN. It is further believed that types IFA and IFB (AAN0044, AAN0045) are used as initiators for the preparation of human fibroblast IFN. The transcript derived using IFN is given in Appendix 4. It consists of two inventors claim a multi-class hybrid interferon polypeptide and a DNA unit having a nucleotide sequence which encodes it. First, the AA sequence of alpha and beta interferons (pro-IFN) is (i) the 1-73 AA seq. of Human alpha 1 (and IF2 is the 74-166 AA seq. of Human beta 1) (see AAN0055, AAF1022); or (ii) the 1-41 AA seq. of Human alpha 1 (and IF2 is the 43-166 AA seq. of Human beta 1) (see AAPI056, AAF1027). After poly(I- β -C) re-anino terminal end of a

M-31: US-39 82-694-7.tif (1971) * .tiff format .tif
Date: Oct 9, 2002 3:15 PM
About: Results were produced by the GenoGenie software, version 4.5,
Running on Macintosh Computer.

LITERATURE 1993-2000

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#combined_fasta_fasta.ses
-MODEL=1-frame -IUP2P-model -DEV x1h
-UU -PAI -PSI -QSMI -FASTA -SPLITX=2 -TUP=3 -GAP=0.02505 -GAP3=0.21445 -IUP=3/4/5/6 Fasta_1.144
-DR_PIR=T -QSMI_FASTA -SPLITX=2 -TUP=3 -GAP=0.02505 -GAP3=0.21445 -IUP=3/4/5/6 Fasta_1.144
-GAPEXT=1.000 -MINMAX=0.100 -IUPHET=0.000 -LOOPEXT=0.000
-QSMI=3 -GAP=0.02505 -GAP3=0.21445 -NCALIB=100000 -XTRANEXT=0.500
-GAPIN=0.000 -GAPMX=1.7 -TUP=3 -YCALIBP=100000 -YGAPXP=0.500
-DELIN=6.000 -DELETE=7.000 -START=1 -MATRIX=biomes62
-TRANS=human4.0.edb -LIST=4.5 -DOCALIGN=200 -THR_SQPF_PSI
-THR_MAX=1.00 -THR_MIN=0 -ALIGN=1.5 -MOLDE=LOCAL -OUTFILE=PTIS
-PER_EX=1 -HEATMAP=500 -MINLEN=0 -MAXLEN=2000000000
-NCALIB=1000000000 -NCPD=6 -IUPHET=3 -IUPLOG
-NCALIB=1000000000 -NCPD=6 -IUPHET=3 -IUPLOG
-WARN_LIMED=0.3 -NO_XLIPXY -WAIT -THREADS=1

Scarf -h information block:
Scarf -h 0.8.3.2 -658 7
Query -Voronoi: 8
Database: PIR 7.1; *
Database sequences: 283138
Database: seqdb: 96009334
Database: seqdb: 23,450000
Scarf -h time (sec): 23,450000

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... 2330
04479; 15611^t;
ian, R.; Gedde

seq_name: Pnt2_Alt3568
seq_documentation_block:
avirilavin resistance protein F [imported] - *Brucella melitensis* (strain 1448)

Species *Brucella melitensis*, received on 01-Feb-2002 at text change 15 Feb 2002
date 01-Feb-2002 **reference** 01-Feb-2002 **text change** 15 Feb 2002
Accession AB5568 **host** *V. vulnificus* **isolate** B-1004 **major** C-1004 **type** 1

Second, the central bank's independence is greatest if it is not part of the government. This is why the Bank of England is independent while the Reserve Bank of New Zealand is not.

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A:Accession #: AH568
A:Status: Preliminary
A:Material Type: DNA
A:Residence: 1-1025 - KUR

A. A suggestion from 2.1
A. M. consider effects of RNA
A. Posterior distribution
A. An act reflects on 10.40. An asset has
investment. This portion is a good market for writings. timer, and it plays an important role
in portfolio.
A. Index: Dow/Jones
A. M. put position: 114.00
A. Know what you want

Implement suggestions:

1. 1.7

2. 0.0

3. Implement: 4.2.05^{b9}

4. Implement: 1.0

5. Implement: 4.2.05^{b9}

6. Implement: 1.0

7. Implement: 1.0

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99. Implement: 1.0

100. Implement: 1.0

- RX MEDLINE:9154672; PubMed:8589678;
RA SAVITSKY K., SIEV Y., HANNAH J., ZIV Y., SARTIET A., COLLINS F.S.,
RA SHILLOH Y., ROTMAN G.: "The complete sequence of the coding region of the ATM gene reveals
RA similarity to cell cycle regulators in different species.";
RA Hum. Mol. Genet. 4:2025-2032(1995).
RN [42]
RP SPOUSENCE FROM N.A.
RX MEDLINE:97443327; PubMed:9199327;
RA PLATZ M., ROTMAN G., BAKER D.J., UZCIEL T., SAVITSKY K., BARI-SHIRE A.,
RA GIARD S., SHILLOH Y., ROSENTHAL A.;
RA "Ataxiatelangiectasia: longest sequence analysis of 184 kb of human
RNA genomic DNA containing the entire ATM gene.";
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RP SPOUSENCE OF 1-24 FROM N.A.
RX MEDLINE:9726790; PubMed:9199447;
RA SAVITSKY K., PLATZ M., ROTMAN G., BAKER D.J., UZCIEL T., GIARD S., SARTIET A., ROSENTHAL A.,
RA "Ataxiatelangiectasia: structural diversity of untranslated sequences
RA suggests complex post-translational regulation of ATM gene."
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RX MEDLINE:96181449; PubMed:8789452;
RA BYRD P.J., MCCONVILLE C.H., CHAUVEZ E., PARKHILL J., STARKEY T.,
RA MUSCATRE G.M., THICK J.R., TAYLOR A.M.R.;
RA "Mutant mice carrying the serine/proline-rich half of the gene for ataxia-
RA telangiectasia.";
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RN SEQUENCING P-1349-3056 FROM N.A.; AND VARIANT ASN-3003.
RX MEDLINE:961805020; PubMed:8524922;
RA BASTIOLI M., COOKE M.;
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RA telangiectasia.";
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RP SEQUENCE OF 1349-3056 FROM N.A.; AND VARIANT ASN-3003.
DEI: 2546-SER-TLE 2548 DEL AND SER-2860 DEL.
RP TISSUE/Fibroblast:
RX MEDLINE:95312868; PubMed:7792600;
RA SAVITSKY K., BAKER D.J., GIARD S., ROTMAN G., ZIV Y., VANAQUAITE L.,
RA TADDEI B.A., SMITH S., UZCIEL T., SIEV Y., ASHKENAZI M., PECKER I.,
RA FREDMAN M., HARRISON J., FAJUNTAI S.R., SIMPSON A., CINNES G.A.,
RA SARTIET A., GATTI R.A., CHESSA L., SANAI O., LAVIN M.F.,
RA JASPER'S N.G.J., TAYLOR A.M.R., ARLETT C.F., MIKI T., WEISSMAN S.M.,
RA LOVETT M., COLLINS F.S., SHILLOH Y.;
RA "A simian ataxia telangiectasia gene with a product similar to PI-3
RA kinase.";
RA Science 268: 1749-1753(1995).
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RP PAR1A: SEQUENCE FROM N.A., AND VARIANT CYS-49; ARG-1054; PHF-1420;
RX MEDLINE:9620739 AND ALA-2897.
RX MEDLINE:96275738; PubMed:8966563;
RA VODOTOVSKY T., RABIN D., LINI L., MONACO G., HAMMARSTROM L.,
RA WEBSTER A.D.B., ZAKHAROK J., BARTONI-BRODNEK C., JAMES M.R.,
RA RUSSO G., CORSE C.M., MEDINA M.,
RA "The ATM gene and susceptibility to breast cancer: analysis of 38
RA breast cancer cases responsive to mutation.";
RA [49]
RN CANCER RES. 56:2725-2732(1996).
RP PURIFIGATION: PUBMED:89694240;
RA "The product of the ATM gene is a 370-kDa nuclear phosphoprotein.";
RA [50]
RN "The product of the ATM gene is a 370-kDa nuclear phosphoprotein.";
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RP SUBCELLULAR LOCALIZATION.
RX MEDLINE:97203148; PubMed:909066;
RA BROWN K.D., ZIV Y., SATHANASIVAM N., CHESSA L., COLLINS F.S.,
RA SHILLOH Y., TADDEI B.A.;
RN [52]
RP "The ataxia-telangiectasia gene product, a constitutively expressed
RX nuclear protein that is not up-regulated following genotoxin damage.";
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RP FROM: NATL. ACAD. SCI. U.S.A. 94:1840-1845(1997).
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RP SUBCELLULAR LOCATION AND VARIANTS AT 2546-S-1-2548 DEL. AM: Y 29-24.
RX MEDLINE:97295602; PubMed:9150358;
RA WATTERS D., KHANNA K.K., BEAMISH H., BARTOJ G., SPRING K., KEDAR V.,
RA GATEI M., STEINZOGL D., HOLSON K., KUZIOV S., ZHABA N., FARRELL A.,
RA KANSAY J., GATTI R.A., LAVIN M.F.;
RA "Cellular localisation of the ataxia-telangiectasia (ATM) gene product
RT and discrimination between mutated and normal forms.";
RN oncogene 14:1911-1921(1997).
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RP KINASE ACTIVITY: PubMed:8988033;
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RA JUNG B., KOROLEV A., LEE S.A., DIMITCHEV A., DRITSCHKO A.;
RA "ATM gene product phosphorylates κ -kappa B-alpha";
RA Cancer Res. 57:24-27(1997).
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RP C-ARM BINDING: MEDLINE:9711440; PubMed:9168117;
RX SHALMAN T., KHANNA K.K., KEDAR V., SPRING K., KEDAR V.,
RA HOLSON K., GATEI M., ZHABA N., WATTERS D., FECTON M., SHILLOH Y.,
RA KHAIRI S., KUTOK D., LAVIN M.F.;
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RA TAYA Y., GAIN JULI H., CHAN D., LEE MILLER S.P., LAVIN M.F.;
RT "ATM associates with and phosphorylates p53; mapping the region of
RT interaction.";
RN Natl. Cancer. 20: 398-400(1998).
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RP p53-A2P10 B1NDING: MEDLINE:98374320; PubMed:9707615;
RX LIM D.-S., KIRSCH D.G., GAMMON C.E., RHU J.-H., ZIV Y., NEWMAN L.S.,
RA DARNEILI R.B., SHILLOH Y., RASTAN M.B.;
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RA PROG. NATL. ACAD. SCI. U.S.A. 95:10146-10151(1998).
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RP PHOSPHORYLATION OF p53: MEDLINE:9843427; PubMed:9733514;
RX BARNI S., MOYAL T., SHILLOH Y., TAYA Y., ANDERSON C.W., CHESSA L.,
RA SHILLOH-YASHILOVSKY N.I., PRIVES S., REISS Y., SHILLOH Y., ZIV Y.;
RI "enhanced phosphorylation of p53 by ATM in response to DNA damage.";
RA SCIENCE 281:1674-1677(1998).
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RP PHOSPHORYLATION OF p53, AND MUTAGENESIS OF ASN-2870 AND ASN-2875: MEDLINE:98404274; PubMed:9733515;
RX GAMMON C.E., LIM D.-S., CHAPRICH K.A., TAYA Y., LAVIN M.D.;
RA SAKURUCHI K., APPELLA E., KASTAN M.B., SILICATO J.D.;
RA "Activation of ATM kinase by ionizing radiation and
RT phosphorylation of p53.";
RN SCIENCE 281:1677-1679(1998).
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RX JACOBSON S.P.; MEDLINE:9932198; PubMed:10500142;
RT SMITH S.G.H., GARY B.B., LAVIN M.D., HANN B.C., YOO S. H., CHEN D.-J.,
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RX CORTEZ D., WANG Y., OIN J., ELLIOTT S.J.;
RT "Requirement of ATM-dependent phosphorylation of brca1 in the DNA
RT damage response to double-strand breaks.";
RN SCIENCE 284:1152-1166(1999).
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OC
 OX
 RN
 RP
 SEQUENCE FROM N_A;
 RX
 MINIRIC: 99105912; pubmed: 9887327;
 RA
 Borsig G., Deitrich A., Bellalou A., Bulfone A., Bernard L.,
 RA
 Bardi S., Gattiuso C., Martini M., Poxon M., Ponnai D., Morellet R.,
 RA
 Wichterle J., Reale S., Strobl J., Kress J., Kell W., "can recognition via
 RA
 Hanson E.; "EYA", a novel vertebrate gene related to the *moscophora* genes, absent in
 RA
 trout, mouse, zebrafish, 8:11-23 (1999).
 RC
 "EYA", a novel vertebrate gene related to the *moscophora* genes, absent in
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 trout, mouse, zebrafish, 8:11-23 (1999).
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 1 SIMILARITY) MAY BE IN THE EYA FAMILY.
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 DR
 EMBL: AJ007998; CA007821; 1;
 DR
 EMBL: AJ007999; CA007821; 1; JOINED;
 DR
 EMBL: AJ008000; CA007821; 1; JOINED;
 KW
 developmental protein; multi-isopeptide family;
 FT
 N-TER 119 AA: 1-349 MW: 988361133729904 C8064;
 FT
 N-TER 119 AA: 1-349 MW: 988361133729904 C8064;
 SQ
 SEQUENCE

Date: Oct 9, 2012 3:16 PM
 About : Results were produced by the GenoScope software, version 4.0
 Copyright (C) 1993-2010 Computer Ltd.
 out_format : pif

Date: Oct 9, 2012 9:16 PM
 About: Results were produced by the GenoScope software, version 4.5,
 Copyright (c) 1993-2010 Computer Ltd.

us-09-832-658-7.n2p.rspt

OS Saccharum robustum.
 OC Fakarava; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 OS Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACC clade;
 OC Panicoideae; Andropogoneae;
 OX NCBI_TaxID:62334;

[1] RN
 RP SEQUENCE FROM N. A.
 RC TISSUE-STEM APEX;
 RA SPANN-CV. MOLOKAI 5829; TISSUE-STEM APEX;
 RT "Differential expression of soluble acid invertase (SAI) genes
 correlates to differences in sucrose accumulation in sugarcane.";
 RL Submitted (MAY 1998) to the EMBL/GenBank/DDBJ databases.
 CC -1- CATALYTIC ACTIVITY: hydrolases of terminal N-Acetylglucosamine
 CC FRUCTOFURANOSIDE RESIDUES IN BETA-D-FRUCTOFURANOSIDES;
 CC -1- SIMILARITY: BELONGS TO FAMILY 32 OF GLYCOSYL HYDROLASES;
 DR EMBL:AF062734; AAC16545.1;
 DR InterPro:IPR001462; GlycC hydro_32;
 DR IPR00251; Glyco_hydro_32_1;
 DR PROSITE: PS00609; GLYCOSYLY_HYDROL_E32_1;
 KW Glycoprotein; Glycosidase; Hydrolase;
 FT NON-TER 567 AA; 626.51 MW; 42FE2E;18721UB3 CRC64;
 SQ SEQUENCE 567 AA; 626.51 MW; 42FE2E;18721UB3 CRC64;

SRP DOCUMENTATION_BLOCK;
 ID 017729; PRELIMINARY; PRI; 477 AA.
 AT 01-JAN-1998 (REMBL); 05, (created)
 LT 01-JUL-2001 (REMBL); 19, (last annotation update)
 LE 01086; 6; PROTEIN
 RN Blobs, etc.
 OS Cephalothrix elegans.
 RA Eukaryota; Metazoa; Nemataoda; Chironomida; Rhabditida; Rhabditoidae;
 RL Rhabditidae; Psiloderaida; Caenorhabditis;
 RX NCBI_TAXID:6239;

RP SEQUENCE FROM N. A.
 RA SPANN-CV. MOLOKAI 5829; TISSUE-STEM APEX;
 RL Submitted (MAY 1998) to the EMBL/GenBank/DDBJ databases.
 RC SPANN-CV N. A.
 RR MOLOKAI 5829; PubMed:9851916;

RA NUMBER:
 RL "Sequence similarity of the nematode C. elegans: A platform for
 life sciences biology";
 DR SCOPUS:Z8222012201983997;
 DR EMBL:Z01491; SAK402321;
 SEQUENCE 477 AA; 54218 MW; 826 AF004061A231C CRC64;

ALIGNMENT_SCORES:
 QUALITY: 50.00 LENGTH: 17
 GAPS: 0
 PERCENT SIMILARITY: 58.824 PERCENT IDENTITY: 58.824

ALIGNMENT_BLOCK;
 ID 017729; PRELIMINARY; PRI; 477 AA.
 AT 01-JAN-1998 (REMBL); 07, (created)
 LT 01-JUL-2001 (REMBL); 19, (last annotation update)
 LE 01086; 6; PROTEIN
 RN Blobs, etc.
 OS C. elegans

seq_name: sp-plant:065342
 seq_documentation_block;
 ID 065342; PRELIMINARY; PRI; 567 AA.
 AC 065342;
 DT 01-AUG-1998 (REMBL); 07, (created)
 ID 01-AUG-1998 (REMBL); 07, (last sequence update)
 DR SUBSTRATE: ACTIVIN INVERTASE (TREATMENT).
 OS Saccharum officinarum (Saccharum).
 OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 OC Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACC clade;
 OC Panicoideae; Andropogoneae;
 OX NCBI_TAXID:4547;

[1] RN
 RP SEQUENCE FROM N. A.
 RC STRAIN-CV. LOUISIANA PURPLE; TISSUE-STEM APEX;
 RA Albert H.H., Zhu Y.J., Moore P.H.;
 RT "Differential expression of soluble acid invertase (SAI) genes
 correlates to differences in sucrose accumulation in sugarcane.";
 RL Submitted (MAY 1998) to the EMBL/GenBank/DDBJ databases.
 CC -1- CATALYTIC ACTIVITY: HYDROLYSIS OF FRUCTOFURANOSIDES;
 CC -1- SIMILARITY: BELONGS TO FAMILY 32 OF GLYCOSYL HYDROLASES;
 DR EMBL:AF062733; AAC16557.1;
 DR IPR001462; Glyco_hydro_32;
 DR PROSITE: PS00609; GLYCOSYL_HYDROL_E32_1;
 KW Glycoprotein; Glycosidase;
 FT NON-TER 567 AA; 626.47 MW; 47AF084CFB2AB CRC64;
 DR SPANN-CV INVERTASE (FRUITWATER);
 DE

RE: [REDACTED] (FRAGMENT)

Kartikai; Karriaciers (Rat); Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Rodentia; Muridae; Murinae; Rattus.

RX
NBLAXID 1016;
RN
[REDACTED]

SEQUENCE FROM N.A.

KA
Kuro T., Koseki-Hirai K., Xue Y., Shirao T.;
study of processes formation in dermin cDNA transfecting fibroblast

RJ
cells";
RJ
submitted; "NAN..."; "The gene, "nanog", trop, fibroblasts".

RJ
EMBL; ABO1_042; BRAZB746_17_1;

DR
Later PROJ: IFPRO2108; Contamin_ABP:

DR
Pran; PEPR_41; contam_ABP; 1;

DR
SMAK1; SMD0102; ABP; 1;

FJ
NQ_N_TER
661 AA: 72682 MW: HF606FC2889983495 CRC64;

alignment_scores:

quality: 49 50

length: 29

ratio: 2.152 Gaps: 1

percent similarity: 79.310 percent identity: 41.379

alignment_blocks:

US_39_W42_658_7_X_170205 ..

Align seq1 /: to: 070205 from: 1 to: 661

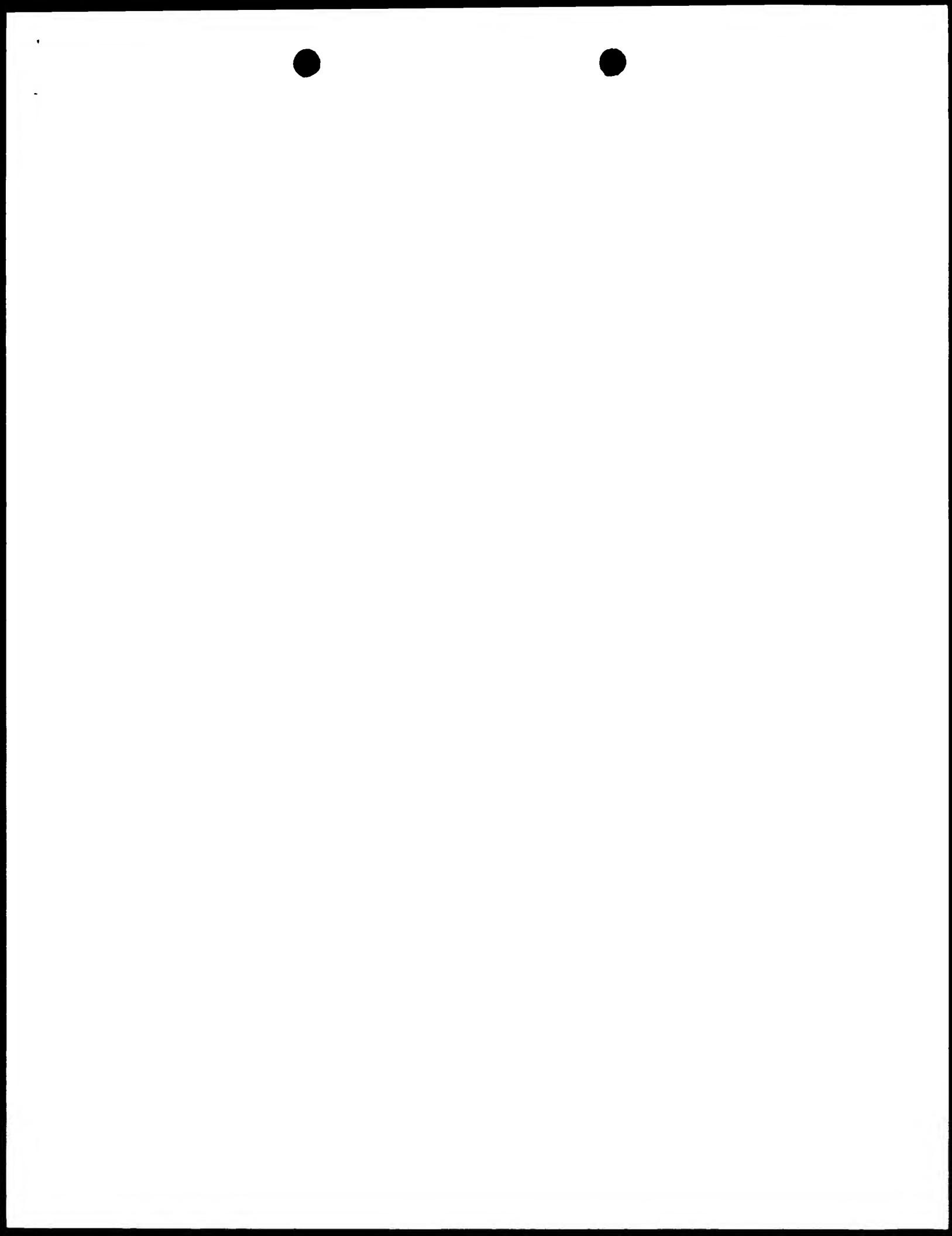
Align seq1 /: to: 070205 from: 1 to: 661

47 GAGAATGAGATGAGAAGTGTACCGGGCTTGAGCCCTCAA.. 50

47 Glyceraldehyde-3-phosphate dehydrogenase; glyceraldehyde-3-phosphate dehydrogenase; G3PDH 53

c1 .GTCCTGAAATTGAGTGCAACGCCCTGG 86

c4 .GTCCTGAAATTGAGTGCAACGCCCTGG 86



Fri Oct 11 10:13:58 2002

us-09-832-658-7.n2P.rai

COMPUTER READABLE FORM
MEDIUM TYPE: FLOPPY DISK

OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PATENT RELEASE #1.0, VERSION #1.25

CURRENT APPLICATION DATA: US-09-832-658

APPLICATION NUMBER: US-09-832-658

FILING DATE: 1999-09-12

CLASSIFICATION:

PRIOR APPLICATION DATA: 09/475,774

FILING DATE: 1998-07-01

REFERENCE/DOCKET NUMBER: 27,794

TELECOMMUNICATION INFORMATION:

NAME: BAILEY, JR., James F.

REGISTRATION NUMBER: 27,794

TELEPHONE: (212) 596-9000

TELEFAX: (212) 596-9090

REF SEQ ID NO: 1;

SEQUENCE CHARACTERISTICS:

LENGTH: 166 amino acids

TYPE: amino acid

STRANGENESS: single

TOPOLOGY: Linear

MOLECULE TYPE: protein

HYPOTHETICAL: NO

ANTI-SENSE: NO

US-09-912-768-1

alignment_scores:
quality: 89.00 length: 22
ratio: 4.944 gaps: 0
Percent Similarity: 81.818 Percent Identity: 77.273

alignment block:
US-09-832-658-7 x US-09-397-992A-7

Alignment seq 1/1 to: US-09-397-992A-7 from: 1 to: 166

seq_documentation block:

Sequence 1, Application PG/TUS9503205

GENERAL INFORMATION:

APPLICANT: Biogen, Inc.

APPLICANT: Gori, Susan E.

APPLICANT: Cate, Richard L.

APPLICANT: Pepinsky, Blake R.

APPLICANT: Chow, Pingchuan E.

TITLE OF INVENTION: Novel Molecules of TEP-Beta

CORRESPONDENCE ADDRESS:

ADDRESSEE: Pish & Neave, 1251 Avenue of the Americas

STREET: New York

CITY: New York

STATE: USA

CITY: 10020-1124
COMPUTER READABLE FORM:
MEDIUM TYPE: FLOPPY DISK
COMPUTER: IBM PC COMPATIBLE
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PATENT RELEASE #1.0, VERSION #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: PG/TUS9503206
FILING DATE:
CLASSIFICATION:
ATTORNEY/AGENT INFORMATION:
NAME: BAILEY, JR., James F.
REGISTRATION NUMBER: 27,794
REFERENCE/DOCKET NUMBER: H179
TELECOMMUNICATION INFORMATION:
TELEPHONE: (212) 596-9000
TELEFAX: (212) 596-9090
INFORMATION SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 166 amino acids
TYPE: amino acid
STRANGENESS: single
TOPOLOGY: Linear
MOLECULE TYPE: protein
HYPOTHETICAL: NO
ANTI-SENSE: NO
PG/TUS9503206-1

alignment_scores:
quality: 89.00 length: 22
Percent Similarity: 81.818 Percent Identity: 77.273

SEQUENCE DATA:
US-09-912-768-1

self-decorationation block:
PATENT NO.: 6326859
APPLICANT: Shudo, Haruo; Muramatsu, Masami; Taniguchi, Tadatosu
TITLE OF INVENTION: DNA AND RIBONUCLEIC PLASMID
NUMBER OF SEQUENCES: 3
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US2001-1459
FILING DATE: 27-OCT-1990
SEQUENCE ID: 1
LENGTH: 187
3'24859-1

